EXHIBIT 4



MANZI MACH 1 - INSTRUMENT CLEANER - PROCESSOR

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1.0 CLASSIFICATION, COMMON OR USUAL NAME, DEVICE NAME

a) Classification Name: Pending - Class II

Common / Usual Name: Endoscope and accessories

Device Classification: 21 CFR § 876.1500, -Endoscope and accessories

Proprietary Name: Manzi Mach 1 Instrument Cleaner- Processor System with MS10 High

Level Disinfector

b) Classification Name: Pending - Class II

Common / Usual Name: Endoscope and accessories

Device Classification: 21 CFR § 876.1500,

Proprietary Name: Manzi MS10

2.0 PREDICATE DEVICE

a) System 83 Plus™ Washer-Disinfector, K983017 Manzi Cleaner System, K043314 (Washer)

b) Steris 20 Sterilant, K875280

3.0 INDICATIONS FOR USE

The Manzi Mach 1 Instrument Cleaner Processor System is indicated for use with the High Level Disinfectant MS10 concentrate (MEC 0.49% PAA, minimum contact temperature of 120°F for a contact time of 15 minutes) for cleaning and high level disinfecting flexible bronchoscopes used in health care settings by health care workers.

4.0 DESCRIPTION OF THE DEVICE

The Manzi Mach 1 Instrument Cleaner-Processor System consists of a Manzi Mach 1 Instrument Cleaner-Processor; a proprietary Manzi germicide, MS10; and a proprietary Manzi Detergent, MD10.

The Manzi Mach 1 Instrument Cleaner-Processor is a self-contained stand-alone system of hardware and software designed to clean and provide high level disinfection of bronchoscopes using the MD10 detergent, the MS10 germicide, and a patented push-pull agitation system. The push-pull agitation system effectively scrubs the interior and exterior surfaces of the bronchoscope without the use of special connectors. The scope is placed in a processing chamber where it is exposed to a push-pull



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agitation cleaning cycle followed by two hot water rinses, a push-pull agitation disinfection cycle that provides high level disinfection (spore log reduction of \geq 10 6 microorganisms with no CFUs) of the device, and a final rinse with an ozonated sanitized water rinse.

The hardware for the Manzi Mach 1 Instrument Cleaner-Processor consists of a stainless steel processing chamber, a push-pull agitation pump, an ozonator, and a variety of components that are mounted in a movable covered frame. The cleaner-processor system utilizes accessories such as disposable water filters, reusable bronchoscope trays, and printer paper.

The Manzi Mach 1 Instrument Cleaner-Processor is designed to: (1) be used in accordance with the reprocessing instructions provided in the operator's manual of the instruments being processed, and (2) facilitate the health care facility's compliance with reprocessing guidelines published by SGNA, APIC, AORN, ASGE, CDC, and other professional organizations.

MD10 is a low foaming enzyme chemical detergent packaged in single use containers for attachment to the Manzi Cleaner. MD10 is intended to be used with the Manzi Instrument Cleaner-Processor.

MS10 is a peracetic acid based liquid chemical germicide. MS10 is intended to be used with the Manzi Instrument Cleaner-Processor.

5.0 SUMMARY OF NONCLINICAL TESTS for the MANZI MACH 1 INSTRUMENT CLEANER-PROCESSOR

5.1 Qualification Testing - FDA Guidance

The Manzi Mach 1 Instrument Cleaner-Processor System was tested and found to conform with the requirements of the "Guidance on Premarket Notification [510(k)] Submissions for Automated Endoscope Washers, Washer / Disinfectors, and Disinfectors Intended for Use in Health Care Facilities", dated August, 1993". The table below identifies the qualifications performed and the results obtained:

Requirement		Requirement	Results
G Performance Data		e Data	
李書聯	1 Process Parameter Tests		Passed
	2 Simul	ated Use Tests	Passed
	2.c. Effe	ectiveness Tests	
	2.c.(1)	Cleaning Efficacy	Passed
	2.c.(2)	Disinfection Efficacy	Passed
	2.c.(3)	Rinsing Efficacy	Passed
	2.c.(4)	Other Tests	Passed
	2.c.(5)	Combined Process	Passed
建 联位		Jse Tests	Passed
H.	Software Do	ocumentation	Passed
I.	Toxicologic	al Evaluation of Residues	Passed



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The Manzi Mach 1 Instrument Cleaner-Processor System was also tested and found to conform with the requirements of the Draft prEN ISO 15883-1: 2003, Washer-disinfectors – Part 1: General Requirements, Definitions and Tests and Draft prEN ISO 15883-4: 2001, Washer-disinfectors – Part 4: Requirements and Tests for Washer-Disinfectors Employing Chemical Disinfection for Thermo-Labile Endoscopes as identified in the Table below.

prEN ISO 15883-1: 2003 Requirement		Results
6.10	Cleaning Efficacy – Scope Ninhydrin Horse serum- prEN 15883-4 Annex B.1.1	Passed
Annex B Annex E	Cleaning Efficacy – Surrogate Ninhydrin	Passed
	Horse serum- prEN 15883-4 Annex B.1.1	
	Surrogate - prEN 15883-4	
6.11	Disinfection Efficacy – Scope	Passed
Annex D	Sheep blood - prEN 15883-4 Annex D	
	Disinfection Efficacy – Surrogate	Passed
	Sheep blood - prEN 15883-4 Annex D	

5.3 Qualification Testing - Langford IC Systems (LIC) Requirements

The Manzi Mach 1 Instrument Cleaner-Processor System was also tested and found to conform to the LIC requirements identified in the table below.

LIC Cleaning Requirement	Results
Cleaning Efficacy: Reduction of protein loading of scopes	Passed
contaminated with a Protein Laden Soil to Remaining Protein	Remaining Protein
levels of $< 6.4 \mu g/cm^2$ (Ref: AAMI TIR30: 2003, A	levels of $< 4.0 \mu g/cm^2$
Compendium of Processes, Materials, Test Methods, and	
Acceptance Criteria for Cleaning Reusable Medical Devices.).	

LIC High Level Disinfection Requirement	Results
High Level Disinfection Efficacy: Reduction of ≥ 10 6 microbial	Passed
loading of scopes with no colony forming units (CFUs)	≥ 6 spore log reduction with no CFUs

LIC Microbiological Efficacy Tests			
Test Method	Test Organisms	Results	
Simulated Use Test (15	Bacillus subtilis	> 6 spore log reduction; no Colony Forming	
min.) ISO 15883-4 Surrogates Sheep's blood soil	Mycobacterium terrae Candida albicans Enterococcus faecium	Units (CFU)	
Sheep's blood son	Styphlococcus aureus		
Simulated Use Test (15 min.) Olympus Bronchoscopes Sheep's blood soil	Bacillus subtilis	> 6 spore log reduction; no CFU	
Simulated Use Test	Candida albicans, Staphylococcus	> 6 spore log reduction;	
Ozonated Water System	aureus, Bacillus subtilis	no CFU	



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LIC Sanitized Ozonated Water System Requirement	Results
Final Rinse System Efficacy: Reduction of ≥ 10 6 microbial	Passed
loading of scopes with no colony forming units (CFUs)	\geq 6 spore log reduction
	with no CFUs

6.0 SUMMARY OF NONCLINICAL TESTS for the MANZI MS10

5.1 Qualification Testing – FDA Guidance

The Manzi MS10 germicide was tested to and met the requirements of the current edition of "Guidance for Industry and FDA Reviewers, Content and Format of Premarket Notification [510(k)] Submissions for Liquid Chemical Sterilants / High Level Disinfectants", dated January 3, 2000. The table below identifies the qualifications performed and the results obtained:

Requirement		Results
5.4	Potency Test	Passed
5.5	Simulated Use Tests	Passed
5.6	In-Use Tests	Passed
6.0	Biocompatibility	Passed

Microbiological Efficacy Summary			
Test Method	Test Organisms	Results	
Sporicidal Activity of Sterilants;	Bacillus subtilis	> 6 spore log reduction;	
AOAC Official Method 966.04	Clostridium sporogenes	No CFUs	
		MS10 is sporicidal	
Fungicidal Activity of Sterilants;	Trichophyton	MS10 is fungicidal	
AOAC Official Method 955.17	mentagrophytes		
Use-Dilution Method; AOAC	Salmonella choleraesuis	MS10 is bactericidal	
Official Method 955.14, 955.15,	Staphylococcus aureus		
964.02	Pseudomonas aeruginosa		
Virucidal Testing	Poliovirus Type 1	MS10 is virucidal	
Quantitative Tuberculocidal Test	Mycobacterium bovis	MS10 is tuberculocidal	

7.0 OVERALL PERFORMANCE CONCLUSIONS

The studies demonstrate that the Manzi Mach 1 Instrument Cleaner-Processor System is safe and effective for the cleaning and high level disinfection of bronchoscopes within the stated indications for use for the Manzi Mach 1 Instrument Cleaner-Processor, the Manzi MS10 germicide, and the Manzi Detergent, MD10, and establishes substantial equivalence of the Manzi Mach 1 Instrument Cleaner-Processor System to the predicate devices identified in Section 2.0 above.